## Amendments to the Claims

This listing of amended claims will replace all prior versions, and listings, of claims in the specification:

1. (previously amended) A ribonucleoside-derivative of the formula

## wherein

R<sub>1</sub> is a base of the purine- or pyrimidine- family,

R<sub>2</sub> is a proton or a substituted derivative of phosphonic acid,

R<sub>3</sub> is a proton or a protection-group for the oxygen atom in 5'-position,

 $R_4$ ,  $R_5$  and  $R_6$  are independently alkyl, aryl, or heteroatom substituted with 1-3 substituents independently selected from alkyl, aryl, alkyl-aryl or aryl-alkyl, wherein the heteroatom is selected from among O, N, Si, Ge, Sn and Pb; or any two of  $R_4$ ,  $R_5$  and  $R_6$  taken in combination with the Si to which they are attached, form a heterocyclic ring; and

wherein at least one of the  $R_4$ ,  $R_5$  or  $R_6$  substituents comprises a tertiary C-atom or a heteroatom that is directly bonded to the Si-atom.

- (previously amended) A ribonucleoside-derivative according to claim 1 wherein the substituent comprising the tertiary C-atom directly bonded to the Si-atom comprises from 4 to 24 C-atoms.
- (previously amended) A ribonucleoside-derivative according to claim 1 wherein the substituent comprising the tertiary C-atom directly bonded to the Si-atom is an alkylsubstituent selected from the group consisting of tert-butyl, tert-pentyl, tert-hexyl, tert-heptyl, tert-octyl, tert-nonyl, tert-decyl, tert-undecyl, tert-dodecyl.
- 4. (previously amended) A ribonucleoside-derivative according to claim 1 wherein the substituent comprising the tertiary C-atom directly bonded to the Si-atom is selected from the group of 1,1-dimethyl ethyl, 1,1-dimethyl-propyl, 1,1-dimethyl-butyl, 1,1-dimethyl-pentyl, 1,1-dimethyl-pentyl, 1,1-dimethyl-pentyl, 1,1,2-trimethyl-butyl, 1,1,2-trimethyl-pentyl, 1,1,2-trimethyl-pentyl, 1,1,2-trimethyl-propyl, 1,1,2-tetramethyl-butyl.

- 5. (previously amended) A ribonucleoside-derivative according to claim 1 wherein one of R<sub>4</sub>, R<sub>5</sub> and R<sub>6</sub> is a heteroatom substituted with 1-3 substituents independently selected from alkyl, aryl, alkyl-aryl or aryl-alkyl, and wherein the heteroatom is selected from among O, N, Si, Ge, Sn and Pb.
- 6. (previously amended) A ribonucleoside-derivative according to claim 5 wherein the substituent directly bonded to the Si-atom comprises a substituted bivalent heteroatom.
- (original) A ribonucleoside-derivative according to claim 6 wherein the heteroatom is oxygen.
- 8. (currently amended) A method for the preparation of a ribonucleoside-derivative according to claim 1, comprising reacting a nucleoside with the formula

where R<sub>1</sub> and R<sub>3</sub> are as defined in claim 1, with a silyloxymethylderivative silyloxymethyl derivative of the formula

wherein Y is a suitable leaving group and wherein R<sub>4</sub>, R<sub>5</sub> and R<sub>6</sub> are as defined in claim 1.

- 9. (original) The method of claim 8 wherein Y is a halogen.
- 10. (previously amended) The method of claim 8 wherein R<sub>4</sub>, R<sub>5</sub> and R<sub>6</sub> together comprise between 3 and 30 carbon atoms.
- 11. (previously amended) The method of claims 8 wherein R<sub>4</sub>, R<sub>5</sub> or R<sub>6</sub> comprise at least one substituted heteroatom directly bonded to the Si atom.
- 12. (original) The method of claim 11 wherein the heteroatom is a bivalent atom.
- 13. (original) The method of claim 12 wherein the heteroatom is oxygen.

- 14. (previously amended) The method of claim 11 wherein the ribonucleoside derivative is further substituted on the oxygen in 3'-position with a group comprising of a derivative of phosphonic acid.
- 15. (previously amended) A method for the preparation of a ribonucleoside-derivative, comprising reacting a ribonucleoside derivative with the formula

upon an electrophilic activation with a compound of formula:

wherein  $R_1$ ,  $R_4$ ,  $R_5$  and  $R_6$  are defined as in claim 1 and  $R_7$  is a alkyl- or aryl-group, or alkyl-aryl-group,

wherein  $R_2$  is a protecting group, and wherein  $R_3$  is a protecting group.

- 16. (cancelled)
- 17. (previously amended) The method of claim 15 wherein the ribonucleoside derivative is further substituted on the oxygen in 3'-position with a group comprising of a derivative of phosphonic acid.